

PCTWORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C09J 175/04	A1	(11) International Publication Number: WO 98/20087 (43) International Publication Date: 14 May 1998 (14.05.98)
<p>(21) International Application Number: PCT/US97/18391</p> <p>(22) International Filing Date: 8 October 1997 (08.10.97)</p> <p>(30) Priority Data: 08/745,364 8 November 1996 (08.11.96) US</p> <p>(71) Applicant: MINNESOTA MINING AND MANUFACTURING COMPANY [US/US]; 3M Center, P.O. Box 33427, Saint Paul, MN 55133-3427 (US).</p> <p>(72) Inventor: SHIMIZU, Yoshinori; P.O. Box 33427, Saint Paul, MN 55133-3427 (US).</p> <p>(74) Agents: MCGEEHAN, Lisa, M. et al.; Minnesota Mining and Manufacturing Company, Office of Intellectual Property Counsel, P.O. Box 33427, Saint Paul, MN 55133-3427 (US).</p>		<p>(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).</p> <p>Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i></p> <p><i>07 2194</i></p>
<p>(54) Title: MOISTURE-CURABLE, HOT-MELT COMPOSITION</p> <p>(57) Abstract</p> <p>There are provided moisture-curable hot-melt compositions comprising a polyurethane prepolymer and a crystalline polymer obtained from monomer(s) containing at least one crystalline ethylenically unsaturated monomer with no active hydrogen, which have low melt viscosities and high initial impact strength and final strength. The monomers may also contain amorphous ethylenically unsaturated monomers with no active hydrogen, in addition to the crystalline monomer. Typical crystalline monomers are (meth)acrylate esters of non-tertiary alcohols, the alkyl portions of which comprise 16 or more carbon atoms. Typical amorphous monomers are (meth)acrylate esters of non-tertiary alcohols, the alkyl portions of which comprise no more than 15 carbon atoms. Use of an alkyl acrylamide as one of the amorphous monomers is advantageous for improving the performance of the moisture-curable hot-melt adhesive.</p>		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

MOISTURE-CURABLE, HOT-MELT COMPOSITION

Field of the Invention

The present invention relates to a moisture-curable, hot-melt composition.

5 More specifically, the present invention relates to a composition comprising a polyurethane prepolymer modified with a crystalline polymer obtained by polymerizing a crystalline monomer.

Background of the Invention

10 Hot-melt adhesives are solid at room temperature and applied in a molten state to a substrate. The substrate is then mated with another substrate while the adhesive is hot viscous liquid. Upon cooling, it solidifies and exhibits instant strength. Because of this rapid adhesion and easy handling, hot-melt adhesives are widely used in various industries. However, since all fundamental resin components

15 of such hot-melt adhesives are thermoplastic resins, they have the inherent drawbacks of low adhesive strength and poor heat resistance. Consequently, the uses of hot-melt adhesives are presently limited to fields such as packaging and bookbinding which do not require heat resistance and high adhesive strength.

Moisture-curable hot-melt adhesives have been developed for the purpose of

20 overcoming these drawbacks of hot-melt adhesives. These adhesives cure after application by action of moisture.

While conventional hot-melt adhesives are mixtures of a thermoplastic resin such as ethylene-vinyl acetate copolymer (EVA) and a tackifying resin, moisture-curable hot-melt adhesives usually have a polyurethane prepolymer with terminal

25 isocyanate groups as the main component for rapid moisture curing and adhesion after curing, and thus despite being hot-melt adhesives, they are chemically different from conventional hot-melt adhesives, and their performance also differs considerably. For example, although with conventional hot-melt adhesives the strength of the material once it is cooled after application constitutes the final

30 strength, moisture-curable hot-melt adhesives acquire gradually increasing strength after application as the reaction with moisture in the atmosphere proceeds.

Consequently, while the final shear strength of conventional hot-melt adhesives is about the same as the initial strength, moisture-curable hot-melt adhesives usually become fully cured in 5-7 days, reaching a final shear strength of several times higher than those of conventional hot-melt adhesives. However, since the major
5 components of moisture-curable hot-melt adhesives are polyurethane prepolymers with a molecular weight of about a few thousand to a few tens of thousand, a poor initial strength is exhibited immediately after cooling following application, compared to conventional hot-melt adhesives which use thermoplastic resins with a molecular weight of a few hundred thousand. Therefore, it is necessary to
10 temporarily clamp the adhesion site with a clip or the like until the moisture curing progresses to exhibit some degree of material strength, and such use has been difficult in cases where external stress is applied immediately after application of the adhesive.

Many measures have been undertaken to produce moisture-curable hot-melt
15 adhesives with high initial strength including, for example, (1) addition of a tackifying resin (Japanese Unexamined Patent Publication No. 3-88886, etc.), (2) addition of a thermoplastic resin (U.S. Patent Specification Nos. 4,775,719, 4,820,368, 4,808,255, 4,889,915, 5,021,507, 5,418,288, etc.), (3) combination of different polyurethane prepolymers (or polyols) (U.S. Patent Specification Nos.
20 5,173,538, 4,999,407, 4,661,542, 5,115,073, 5,034,453, 4,985,535, etc.) and (4) utilizing a polyurethane prepolymer with a specific structure (U.S. Patent Specification No. 5,019,638, etc.).

Japanese Unexamined Patent Publication No. 3-88886 describes a mixture of a polyurethane prepolymer and a cumarone/indene-based tackifying resin.
25 Though this tackifying resin has the effect of increasing the initial adhesive strength, the heat resistance of the adhesive after curing is low, and problems such as phase separation and coloring also occur by aging.

U.S. Patent No. 4,775,719 describes a mixture of an ethylene-vinyl acetate copolymer (EVA) or ethylene-ethyl acetate copolymer (EEA) and a polyurethane
30 prepolymer. U.S. Patent No. 4,820,368 describes a mixture of styrene-based block copolymer or EVA with a polyurethane prepolymer. U.S. Patent Specification No.

4,808,255 describes a mixture of EVA and a polyurethane prepolymer. U.S. Patent Specification No. 4,889,915 describes a mixture of a thermoplastic polyurethane polymer and a polyurethane prepolymer. This addition of a thermoplastic resin tends to drastically increase the melt viscosity of the moisture-curable hot-melt adhesive. Also, when such thermoplastic resins are added a tackifying agent may also be added to bring out the initial adhesion, but since tackifying agents cause the problems described above, they are not preferred for moisture-curable hot-melt adhesives.

U.S. Patent Specification Nos. 5,173,538 and 4,999,407 describe mixtures of polyurethane prepolymers with a glass transition temperature (T_g) of room temperature or above and polyurethane prepolymers with a glass transition temperature of room temperature or below. U.S. Patent Specification No. 4,661,542 describes a mixture of a linear polyurethane prepolymer and a branched polyurethane prepolymer. U.S. Patent Specification Nos. 5,034,453 and 4,985,535 describe polyurethane prepolymers created based on polyester polyols with different molecular weights. All of the moisture-curable hot-melt adhesives produced by these methods have the drawback of low initial cohesion of the adhesives even though some degree of initial adhesion is achieved.

U.S. Patent Specification No. 5,115,073 describes a polyurethane prepolymer based on a polyester polyol with a high melting point (65°C - 150°C) and a polyester polyol with a relatively low melting point (30°C - 60°C). The adhesive obtained by this method has the disadvantage of being brittle prior to curing.

U.S. Patent Specification No. 5,019,638 describes an aliphatic polyurethane prepolymer with at least 12 methylene groups. The adhesive obtained by this method is also brittle prior to curing as is the one disclosed in U.S. Patent Specification No. 5,115,073, and its initial strength is inadequate.

U.S. Patent Specification No. 5,021,507 describes a mixture of a low molecular weight (10,000-30,000) acrylic resin with no active hydrogen and a polyurethane prepolymer. The adhesive obtained by this method has improved cohesive and adhesive strength, and is also stated as having excellent initial adhesion. The acrylic resin used is believed to be amorphous.

The expanding fields of use for moisture-curable hot-melt adhesives with improved performance over conventional hot-melt adhesives has led to greater expectations for moisture-curable hot-melt adhesives with both low viscosity for easier dispensing and strong initial cohesion. Such moisture-curable hot-melt adhesives must naturally have adequate final strength.

The low viscosity referred to here is a low viscosity upon melting, and specifically it refers to a Brookfield viscosity of 50,000 cP or lower, and preferably 20,000 cP or lower, at 121°C (250°F). The initial cohesion is measured by the rate of crystallization or by an initial creep test, but even when these values are excellent it is often impractical for actual use. This is because the measurements are only measurements of the initial "static" strength, and do not take into consideration the various external stresses exerted after bonding and prior to use when put into actual use. Consequently, the initial cohesion referred to here represents the "dynamic" impact strength. A strong initial cohesive force is, specifically, an impact adhesive strength one minute after adhesion of at least 5 kg·cm, and preferably at least 10 kg·cm according to JIS K6855 (or ASTM D950). Also, the final strength refers to the overlap shear adhesive strength measured according to JIS K6851, and an adequate final strength is a shear adhesive strength of at least 15 kg/cm², and preferably at least 30 kg/cm². The desired characteristics for moisture-curable hot-melt adhesives are summarized in the following table.

	Allowed value	Target value
Melt viscosity (cP, at 120°C)	≤50,000	≤20,000
Initial impact strength (kg·cm)	≥5	≥10
Final strength (overlap shear strength) (kg/cm ²)	≥15	≥30

Summary of the Invention

As a result of much research on the subject explained above, the present inventor has found that addition of a crystalline polymer to a polyurethane prepolymer gives

a moisture-curable hot-melt adhesive with high initial impact strength and final strength, and with a low melt viscosity. The crystalline polymer referred to here is a polymer obtained by polymerization of an ethylenically unsaturated crystalline monomer which has no active hydrogen and therefore is not reactive with the polyurethane prepolymer. A particularly preferred crystalline polymer according to the present invention is a crystalline polymer obtained by polymerization of a crystalline (meth)acrylate ester monomer. As used herein, the term "(meth)acrylate" means methacrylate or acrylate.

Thus, the present invention provides a moisture-curable hot-melt adhesive comprising a polyurethane prepolymer and a crystalline polymer obtained from monomer(s) containing at least one crystalline ethylenically unsaturated monomer with no active hydrogen.

For the purpose of the present invention, the term "crystalline" means that the material has sufficient crystallinity such that it exhibits a measurable melting temperature (T_m), although such a material is not required to be 100% crystalline.

The "crystalline polymer" referred to here is one with a melting point of 15-135°C, preferably 20-120°C and more preferably 25-100°C.

The "crystalline polymer" is a polymer obtained from monomer(s) containing a crystalline monomer at 5-100 wt%, preferably 10-90 wt% and more preferably 30-80 wt%. The weight average molecular weight of the crystalline polymer is generally 5,000-100,000. The weight average molecular weight of the crystalline polymer is preferably greater than 10,000 and up to 100,000, more preferably greater than 30,000 and up to 100,000, and most preferably greater than 30,000 and up to 60,000. Also the crystalline polymer content in the mixture of the crystalline polymer and the polyurethane prepolymer is 5-90 wt%, preferably 5-80 wt% and more preferably 5-70 wt% of the mixture.

Detailed Description of the Invention

The present invention will now be explained in detail. Although the following explanation concerns a moisture-curable hot-melt adhesive, the

composition of the invention may also be effectively employed as a sealing material or coating material.

The moisture-curable hot-melt adhesive of the invention comprises a polyurethane prepolymer and a crystalline polymer obtained from monomer(s)
5 containing at least one crystalline ethylenically unsaturated monomer with no active hydrogen as at least one part of the monomer.

The polyurethane prepolymer is a prepolymer with isocyanate groups on both ends which is obtained by condensation reaction between a polyol component and an isocyanate component, and is one which is commonly used for moisture-
10 curable hot-melt adhesives.

Examples of usable polyols include polyester polyols obtained by ring-opening polymerization of a lactone such as ϵ -caprolactone; polyester polyols obtained from polyhydric alcohols and polybasic acids; polyether polyols such as polyethyleneoxide diol, polypropyleneoxide diol, polybutyleneoxide diol, and
15 polytetramethyleneoxide diol; aliphatic polyols such as hydrogenated polybutadiene diol and ethylenebutylene diol; polybutadiene diol; ethylene oxide addition product of bisphenol A; propylene oxide addition product of bisphenol A; ethylene oxide/propylene oxide addition product of bisphenol A, etc.

Examples of polyhydric alcohols which may be used in polyester polyols
20 obtained from polyhydric alcohols and polybasic acids include ethylene glycol, 1,2-propylene glycol, 1,3-propylene glycol, 1,3-butylene glycol, 1,4-butylene glycol, neopentyl glycol, 1,8-octanediol, diethylene glycol, dipropylene glycol, cyclohexane-1,4-diol, glycerol etc. Examples of polybasic acids which may be used in polyester polyols obtained from polyhydric alcohols and polybasic acids include
25 succinic acid, glutaric acid, adipic acid, pimelic acid, suberic acid, azelaic acid, sebacic acid, decanedioic acid, dodecanedioic acid, phthalic acid, isophthalic acid, terephthalic acid, and trimellitic acid etc.

Examples of polyisocyanates which may be used include diphenylmethane-2,4'-diisocyanate, diphenylmethane-4,4'-diisocyanate, toluene-2,4-diisocyanate,
30 toluene-2,6-diisocyanate, naphthalene-1,5-diisocyanate, triphenylmethane-4,4',4''-triisocyanate, 1,4-phenylene diisocyanate, 4,4'-cyclohexylmethane diisocyanate,

hexamethylene diisocyanate, isophorone diisocyanate, tetramethylxylene diisocyanate, xylene diisocyanate, etc.

A mixture of polyols may be used, and a mixture of polyisocyanates may be used.

5 Various adjuvants or other ingredients may be added to the adhesive composition of the invention to impart or modify particular characteristics. The adjuvants should be added only at a level that does not materially adversely interfere with the adhesion or cause premature curing of the composition. For examples, fillers (e.g., carbon black; fibers; glass, ceramic, metal or plastic bubbles; metal
10 oxides such as zinc oxide; and minerals such as talc, clays, silica, silicates, and the like) tackifiers (non-reactive and/or reactive to polyisocyanates); plasticizers; antioxidants; pigments; UV absorbers; and adhesion promoters, and the like may be added to modify adhesion, strength build-up, tack, flexibility, etc..

Preparation of polyurethane prepolymers is well-known in the technical
15 field, and thus will not be explained in this specification. According to the invention, the adhesive composition may be prepared by adding the isocyanate component after polymerizing the crystalline polymer in the polyol. Alternatively, the crystalline polymer may be mixed with the polyol component and then the isocyanate component added to this mixture to make the adhesive composition.
20 The molecular weight of the polyurethane prepolymer used in the invention may be varied by changing the ratio of isocyanate equivalent to equivalent of total functional groups containing active hydrogens (usually hydroxyl equivalent), with the ratio being generally 1.1 to 2.5. When the ratio is smaller than 1.1, the molecular weight is excessively high, melt viscosity is significantly increased. When
25 the ratio is larger than 2.5, the amount of unreacted isocyanate is excessively high, causing the problem of stability. This ratio (isocyanate equivalent/equivalent of functional groups containing active hydrogen) is preferably 1.2 to 2.2, and more preferably 1.2 to 2.0.

According to the invention, addition of the crystalline polymer to the
30 polyurethane prepolymer in the composition increases the initial impact strength and

final strength of the moisture-curable hot-melt adhesive while also lowering the melt viscosity.

The crystalline polymer is one obtained by polymerization of monomer(s) containing at least one crystalline ethylenically unsaturated monomer with no active
5 hydrogen. The monomers used to make the crystalline polymer may contain, in addition to the crystalline monomer, an amorphous ethylenically unsaturated monomer with no active hydrogen.

The term "amorphous" means that the material does not exhibit a measurable melting point (T_m) although such a material is not required to be 100%
10 amorphous.

The crystalline polymer is defined here as a polymer with a melting point of 15°-135°C. When the melting point of the polymer is below 15°C, sufficient crystallinity at room temperature cannot be obtained. When the melting point of the polymer is above 135°C, the adhesive composition comprising the crystalline
15 polymer and a polyurethane prepolymer needs to be melted at above 135°C in order to get a uniform mixture. This may lead to viscosity increase due to side reaction and/or decomposition of the adhesive composition. The melting point of the crystalline polymer is preferably 20-120°C, and more preferably 25-100°C.

Examples of crystalline ethylenically unsaturated monomers with no active
20 hydrogen include (meth)acrylate esters of nontertiary alcohols, the alkyl portions of which comprise 16 or more carbon atoms, and (meth)acrylate esters of monools containing polyalkylene oxide units, wherein the alkylene portion of the polyalkylene oxide unit contains 2 or more carbon atoms.

Examples of (meth)acrylate esters of non-tertiary alcohols, the alkyl portions
25 of which comprise 16 or more carbon atoms, include, but are not restricted to, hexadecyl (meth)acrylate, octadecyl (meth)acrylate, eicosanyl (meth)acrylate, behenyl acrylate, behenyl methacrylate (for example, BLEMER VMA, product of Nippon Oil & Fats Co.), hexacosanyl (meth)acrylate, UNILIN X5100 (acrylate ester of alkyl alcohol, the alkyl portion of which comprises 28-35 carbon atoms,
30 product of Petrolite Corp.), UNILIN X-5112 (acrylate ester of alkyl alcohol, the alkyl portion of which comprises 36-44 carbon atoms, product of Petrolite Corp.

and UNILIN X-5113 (acrylate ester of alkyl alcohol, the alkyl portion of which comprises 46-53 carbon atoms, product of Petrolite Corp.).

Examples of (meth)acrylate esters of monools containing polyalkylene oxide units, wherein the alkylene portion of the polyalkylene oxide unit contains 2 or more carbon atoms, include, but are not restricted to, methoxypolyethylene glycol methacrylate (for example, NK ESTER M-230G, product of ShinNakamura Chemical Industries), UNITHOX X-5141 (Petrolite Corp.) and UNITHOX X-5142 (Petrolite Corp.). UNITHOX X-5141 and UNITHOX X-5142 are acrylic acid esters containing a polyethylene glycol portion and a polymethylene portion, and are generally represented by the structural formula:

$\text{CH}_2=\text{CH}-\text{COO}-(\text{CH}_2\text{CH}_2\text{O})_x-(\text{CH}_2)_y-\text{CH}_3$, wherein $1 \leq x \leq 50$, $27 \leq y \leq 53$.

A combination of different crystalline monomers may be used.

Amorphous monomers which may be used together with the crystalline monomer to form the crystalline polymer are ethylenically unsaturated monomers with no active hydrogen. More specifically, the amorphous monomers to be used according to the invention include any (meth)acrylate esters except for (meth)acrylate esters of non-tertiary alcohols, the alkyl portions of which comprise 16 or more carbon atoms (i.e. crystalline monomers) (in other words, (meth)acrylate esters of non-tertiary alcohols, the alkyl portions of which comprise no more than 15 carbon atoms, as well as (meth)acrylate esters of any alcohol other than alkyl alcohols (in which case the number of carbon atoms of the alcohol is not limited), and further include vinyl esters, vinyl ethers, alkyl acrylamides, styrene, acrylonitrile, N-vinyl lactams, vinylidene chloride, vinyltoluene, etc.

Examples of amorphous (meth)acrylate esters include (meth)acrylate esters of alkyl alcohols, the alkyl portions of which comprise no more than 15 carbon atoms, such as methyl (meth)acrylate, ethyl (meth)acrylate, butyl (meth)acrylate, 2-ethylhexyl (meth)acrylate, isooctyl (meth)acrylate, and lauryl (meth)acrylate, and (meth)acrylate esters of alcohols other than alkyl alcohols such as isobornyl (meth)acrylate, 2-phenoxyethyl (meth)acrylate, tetrahydrofurfuryl (meth)acrylate, abitol acrylate, and dicyclopentadienyl (meth)acrylate, but the amorphous (meth)acrylate esters are not limited to these.

Examples of vinyl esters to be used as amorphous monomers include vinyl propionate, vinyl acetate, vinyl pivalate, vinyl neononanoate, vinyl-2-ethyl hexanoate and vinyl neodecanoate. Examples of alkyl acrylamides include N,N-dimethylacrylamide and N,N-diethylacrylamide. Examples of N-vinyl lactams
5 include N-vinylpyrrolidone and N-vinylcaprolactam.

For amorphous monomers, it is possible to use a combination of different monomers.

According to the invention, it has been found that using a crystalline polymer obtained from a monomer containing an alkylacrylamide such as
10 dimethylacrylamide or diethylacrylamide particularly decreases the melt viscosity of the moisture-curable hot-melt adhesive, while also providing excellent initial impact strength and extremely high final strength after curing. In most cases, the alkyl acrylamide is preferably present in the monomer mixture at 5-70 wt%. If the alkyl acrylamide content is less than 5 wt%, the effects on lowered melt viscosity, and
15 improved initial impact strength and shear strength are small, and the content of greater than 70 wt% causes increased melt viscosity. A more preferred range for the alkyl acrylamide is 5-60 wt%.

The crystalline polymer obtained by polymerization of the crystalline monomer has notably improved compatibility with the polyol in the polyurethane prepolymer. The improved compatibility with the polyol means improved
20 compatibility with the polyurethane prepolymer prepared from the polyol, and thus improvement in the characteristics of the adhesive of the invention may be expected. Also, when the crystalline polymer obtained from the crystalline monomer is mixed with the polyol which is the starting material for the polyurethane prepolymer, the
25 mixture has a lower melt viscosity compared to mixtures of amorphous polymers prepared from amorphous monomers. A lower melt viscosity of the polyol dissolving the polymer means that the melt viscosity of the composition containing the polymer mixed with the prepolymer prepared from that polyol will be lower than the melt viscosity of compositions containing amorphous polymers. The hot-
30 melt adhesive is coated on the adherend while molten and immediately hardens upon cooling, and therefore the lower melt viscosity results in easier penetration of

the adhesive into the fine irregularities on the surface of the adherend; thus, a low melt viscosity is very advantageous for improving the adhesive properties of the hot-melt adhesive.

5 The crystalline monomer may constitute 5-100 wt% of the monomer mixture. When a monomer mixture containing less than 5 wt% of a crystalline monomer is used, the compatibility of the resulting polymer with the polyol for the polyurethane prepolymer is lowered, which means lower compatibility with the polyurethane prepolymer itself. According to the invention, the monomer mixture may contain 100 wt% crystalline monomer(s), although the adhesive tends to be
10 brittle in such cases. Consequently, depending on the use, the polymer used is preferably prepared from a monomer mixture containing an amorphous monomer in addition to the crystalline monomer. The crystalline monomer content of the monomer starting substance is preferably 10-90 wt%, and more preferably 30-80 wt%.

15 The weight average molecular weight of the crystalline polymer may be from 5,000 to 100,000. Crystalline polymers with weight average molecular weights of less than 5,000 provide less initial impact strength for the moisture-curable hot-melt adhesives to which they are added. When the weight average molecule of the crystalline polymer is greater than 100,000 the melt viscosity is
20 increased, thus impairing the usefulness of the adhesive. The crystalline polymer preferably has a weight average molecular weight of greater than 10,000 and up to 100,000, more preferably greater than 30,000 and up to 100,000, and most preferably greater than 30,000 and up to 60,000.

The amount of the crystalline polymer present in the mixture of the
25 crystalline polymer and a polyurethane prepolymer may be 5-90 wt% when the crystalline polymer content is less than 5 wt%, less initial impact strength is provided for the moisturecurable hot-melt adhesive. When the crystalline polymer content is greater than 90 wt%, the heat resistance of the adhesive after curing is lowered. The initial impact strength, final strength and melt viscosity of the
30 moisture-curable hot-melt adhesive of the invention appear to be more greatly balanced when the crystalline polymer content is at the lower end of the ranges

specified above. A preferred crystalline polymer content is 5-80 wt%, with 5-70 wt% being more preferred.

The crystalline polymer to be used according to the invention may be synthesized by radical, anionic or cationic polymerization of the monomers comprising the crystalline monomer and the amorphous monomer, although
5 synthesis by radical polymerization is preferred for ease of reaction with a greater variety of usable monomers. The initiator for the radical polymerization may be a thermal initiator which generates radicals by heat, or a photoinitiator which generates radicals by light.

10 Examples of thermal initiators which may be used include azo compounds such as 2,2'-azobis(2,4-dimethylvaleronitrile), 2,2'-azobisisobutyronitrile, 2,2'-azobis(2-methylbutyronitrile), 1,1'-azobis(1-cyclohexane-1-carbonylnitrile) and dimethyl-2,2'-azoisobutyrate, as well as peroxides such as benzoyl peroxide, lauroyl peroxide and t-butyl peroxyphthalate.

15 Examples of photoinitiators which may be used include benzoin ethers such as benzoin methyl ether and benzoin butyl ether, acetophenone derivatives such as 2,2-dimethoxy-2-phenylacetophenone and 2,2-diethoxyacetophenone, and acylphosphine oxide and acylphosphonate derivatives such as diphenyl-2,4,6-trimethylbenzoylphosphine oxide, isopropoxy(phenyl)-2,4,6-trimethylbenzoylphosphine oxide and dimethylpivaloylphosphonate.
20

A chain transfer agent may also be used during synthesis of the crystalline polymer to adjust the polymer molecular weight. Chain transfer agents which may be used are mercapto compounds such as dodecylmercaptan and halogen compounds such as carbon tetrabromide.

25 The moisture-curable hot-melt adhesive of the invention may be easily produced by combination of various well-known techniques. For example, a moisture-curable hot-melt adhesive invention may be produced by first preparing the crystalline polymer, mixing the polymer with the polyol component which is to serve as the polyurethane prepolymer raw material, and then adding an isocyanate
30 component to the mixture for reaction with the polyol component. Alternatively, a moisture-curable hot-melt adhesive according to the invention may be produced by

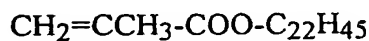
polymerizing the starting monomer in the polyol component which is to be the polyurethane prepolymer raw material to synthesize the crystalline polymer, and then adding an isocyanate component to the mixture for reaction with the polyol to make the polyurethane prepolymer. Yet another alternative method of producing a moisture-curable hot-melt adhesive according to the invention involves preparing the crystalline polymer and polyurethane prepolymer each separately, and then mixing them together. A few of these methods are explained by way of the following examples.

10 Examples

The present invention will now be explained in further detail by way of the following examples. These examples are provided as illustration of the invention and are in no way intended to be restrictive.

The abbreviations used in the examples are the following.

15 VMA: behenyl methacrylate (crystalline monomer, product of Nippon Oil & Pats Co.)



BMA: n-butyl methacrylate (amorphous monomer)

MMA: methyl methacrylate (amorphous monomer)

20 IOA: isooctyl acrylate (amorphous monomer)

DMAA: N,N-dimethylacrylamide (amorphous monomer)

DEAA: N,N-diethylacrylamide (amorphous monomer)

DM: n-dodecylmercaptan

V-65: 2,2'-azobis(2,4-dimethylvaleronitrile) (product of Wako Pure
25 Chemical Industries Co.)

Mw: weight average molecular weight

Mn: number average molecular weight

Mw/Mn: ratio of weight average molecular weight to number average
molecular weight

30 Tm: melting point

SANNIX PP1000: polypropylene glycol with hydroxyl value of 111 (product of Sanyo Chemical Industries)

RUCOFLEX S105-30: Poly(hexamethylene adipate)diol with hydroxyl value of 29.3 (product of Ruco Polymer Co.)

5 MILLIONATE MT-F: 4,4'-diphenylmethane diisocyanate (MDI) (product of Nippon Polyurethane Industries)

DMDEE: dimorpholino diethyl ether (product of Huntsman Co.)

NCO/OH ratio: Equivalent ratio of isocyanate to hydroxyl groups

The various properties of the crystalline polymers were tested in the
10 following manner.

The weight average molecular weight (M_w) and number average molecular weight (M_n) were measured by gel permeation chromatography (GPC).

Tetrahydrofuran was used as the solvent to obtain a molecular weight with polystyrene as the standard. The GPC method is described in detail in Modern Size
15 Exclusion Liquid Chromatography, Practice of Gel Permeation Chromatography, John Wiley & Sons.

The melting point (T_m) was measured using a differential scanning calorimeter (DSC) (PERKIN ELMER 2C) at a temperature-elevating rate of 10°C/min. DSC is described in detail in Thermal Characterization of Polymeric
20 Materials, Academic Press, New York.

In Examples 1-4 described below, the polymers were prepared by the following general preparation of acrylic polymers. Specifically, a monomer (total weight: 30 grams) DM (chain transfer agent) and V-65 (initiator) were placed in a heat-resistant glass bottle (product of Schott Glaswerke) and nitrogen gas was
25 introduced for 15 minutes. The glass bottle was then totally sealed and tumbled in a constant temperature bath (TAIYO Thermo Unit T-368) at 60°C for 4 hours for polymerization.

Example 1

30 This example concerns preparation of amorphous polymers. Amorphous polymers were synthesized using 64.0 parts by weight BMA and 36.0 parts by weight MMA

as amorphous monomers. The amount of the polymerization initiator V-65 was consistently 0.20 parts by weight while the amount of the chain transfer agent DM was varied as shown in Table 1, to obtain amorphous polymers 1-5 having different molecular weights. The molecular weights of the synthesized polymers were then
5 measured. The polymers had no melting point as measured by DSC. The results are summarized in Table 1.

Table I

	Polymer 1	Polymer 2	Polymer 3	Polymer 4	Polymer 5
BMA (monomer)	64.00	64.00	64.00	64.00	64.00
MMA (monomer)	36.00	36.00	36.00	36.00	36.00
DM (chain transfer agent)	0.10	0.30	0.50	1.00	1.50
V-65 (initiator)	0.20	0.20	0.20	0.20	0.20
Polymer type	amorphous	amorphous	amorphous	amorphous	amorphous
M _w	214000	91200	57300	31400	22500
M _n	95000	43200	28200	16000	11600
M _w /M _n	2.25	2.11	2.04	1.96	1.94
T _m (°C)	--	--	--	--	--

Example 2

This example concerns preparation of crystalline polymers.

The general preparation of acrylic polymers described above was used to prepare
5 crystalline polymers from a monomer mixture comprising 50.0 parts by weight
VMA (crystalline monomer), 32.0 parts by weight BMA and 18.0 parts by weight
MMA. The amount of the initiator V-65 was consistently 0.20 parts by weight,
while the amount of DM was varied as shown in Table 2, to synthesize crystalline
polymers 6-11 having different molecular weights. The molecular weights and
10 melting points of the polymers were then measured. The results are summarized in
Table 2.

Table 2

	Polymer 6	Polymer 7	Polymer 8	Polymer 9	Polymer 10	Polymer 11
VMA (crystalline monomer)	50.00	50.00	50.00	50.00	50.00	50.00
BMA (monomer)	32.00	32.00	32.00	32.00	32.00	32.00
MMA (monomer)	18.00	18.00	18.00	18.00	18.00	18.00
DM (chain transfer agent)	0.10	0.30	0.50	1.00	1.50	3.00
V-65 (initiator)	0.20	0.20	0.20	0.20	0.20	0.20
BMA/MMA ratio	64/36	64/36	64/36	64/36	64/36	64/36
Polymer type	crystalline	crystalline	crystalline	crystalline	crystalline	crystalline
M _w	211000	113000	63300	32800	23400	13000
M _n	102000	58000	29700	17300	13320	7600
M _w /M _n	2.07	1.95	2.13	1.90	1.76	1.71
T _m (°C)	34	36	34	39	38	40

Example 3

The general preparation of acrylic polymers described above was also used in this example to prepare crystalline polymers, except that the BMA/MMA ratio of the amorphous monomer was consistently 64/36 as in Examples 1 and 2, while the amount of crystalline monomer VMA was varied, as shown in Table 3. The amounts of DM and V-65 were kept constant. The amounts of each component and the molecular weights and melting points of the resulting crystalline polymers 12-17 are summarized in Table 3.

Table 3

	Polymer 12	Polymer 13	Polymer 14	Polymer 15	Polymer 16	Polymer 17
VMA (crystalline monomer)	80.00	70.00	60.00	50.00	40.00	30.00
BMA (monomer)	12.80	19.20	25.60	32.00	38.00	44.80
MMA (monomer)	7.20	10.80	14.40	18.00	21.60	25.20
DM (chain transfer agent)	1.00	1.00	1.00	1.00	1.00	1.00
V-65 (initiator)	0.20	0.20	0.20	0.20	0.20	0.20
BMA/MMA ratio	64/36	64/36	64/36	64/36	64/36	64/36
Polymer type	crystalline	crystalline	crystalline	crystalline	crystalline	crystalline
M _w	29600	31600	33400	32800	34100	34000
M _n	16700	17800	18500	17300	16500	16300
M _w /M _n	1.77	1.78	1.80	1.90	2.06	2.08
T _m (°C)	53	51	44	39	30	25

Example 4

This example concerns crystalline polymers prepared from monomer starting substances containing alkyl acrylamides.

- 5 Monomer mixture containing DMAA and DEAA as amorphous alkyl acrylamide monomers in the amounts shown in Table 4 were made to prepare crystalline polymers according to the general preparation of acrylic polymers described above. As shown in Table 4, the amount of the polymerization initiator V-65 was kept constant, while the amount of the chain transfer agent DM was
- 10 varied to prepare crystalline polymers 18-23 with different molecular weights. The molecular weights and melting points of the polymers are summarized in Table 4.

Table 4

	Polymer 18	Polymer 19	Polymer 20	Polymer 21	Polymer 22	Polymer 23
VMA (crystalline monomer)	50.00	50.00	50.00	50.00	50.00	50.00
IOA (monomer)	10.25	10.25	10.25	10.25	10.42	10.42
DMMA (monomer)	39.75	39.75	39.75	39.75	0	0
DEAA (monomer)	0	0	0	0	39.58	39.58
DM (chain transfer agent)	1.20	1.60	2.00	2.50	2.00	2.50
V-65 (initiator)	0.20	0.20	0.20	0.20	0.20	0.20
Polymer type	crystalline	crystalline	crystalline	crystalline	crystalline	crystalline
M _w	56600	42700	36400	28700	59900	50700
M _n	15000	12100	10000	8174	9700	8000
M _w /M _n	3.77	3.54	3.64	3.51	6.17	6.34
T _m (°C)	47	46	47	48	45	45

Example 5

This example concerns polymerization of crystalline polymers in a polyol as the polyurethane prepolymer starting material.

- The monomers (total weight: 31.97 grams), SANNIX PP1000
5 (polypropylene glycol, 11.51 grams), DM (chain transfer agent) and V-65 (initiator) were placed in a heat-resistant glass bottle (product of Schott Glaswerke) and nitrogen gas was introduced for 15 minutes. The glass bottle was then totally sealed and tumbled in a constant temperature bath (TAIYO Thermo Unit T-368) at 60°C for 6 hours for polymerization. The compositions of the monomer starting
10 substances and the molecular weights and melting points of the resulting crystalline polymers 24-27 are summarized in Table 5.

Table 5

	Polymer 18	Polymer 19	Polymer 20	Polymer 21
VMA (crystalline monomer)	0.00	50.00	30.00	50.00
BMA (monomer)	64.00	32.00	44.88	0.00
IOA (monomer)	0.00	0.00	0.00	10.25
MMA (monomer)	36.00	18.00	25.20	0.00
DMMA (monomer)	0	0	0	39.75
SANNIX PP1000 (polypropylene glycol)	25.12	25.12	25.12	25.12
DM (chain transfer agent)	1.00	1.00	1.00	1.00
V-65 (initiator)	0.20	0.20	0.20	0.20
Polymer type	amorphous	crystalline	crystalline	crystalline
M_w^*	37100	37100	37300	53200
M_n^*	18300	19400	18900	20200
M_w/M_n	2.02	1.90	1.98	2.64

* Molecular weight of polymer alone (without polypropylene glycol)

Example 6

Some of the amorphous polymers and crystalline polymers synthesized above were evaluated for compatibility with a polyol. The amorphous polymer or crystalline polymer was placed in a small metal container with SANNIX PP1000

- 5 (polypropylene glycol), and the container was heated on a hot plate at 120°C for 2 hours, after which the contents were thoroughly stirred. Samples with weight ratios of the amorphous polymers or crystalline polymers to SANNIX PP1000 of 1/9, 1/3 and 1/1 were each tested. The stirred samples were observed for solubility and uniformity of mixture of the polymers with the polyol. The results are summarized
- 10 in Table 6.

Table 6

Polymer type	Polymer/SANNIX = 1/9	Polymer/SANNIX = 1/3	Polymer/SANNIX = 1/1
Polymer 1 (amorphous polymer) (VMA%=0, Mw=214000)	insoluble, non-uniform	insoluble, non-uniform	insoluble, non-uniform
Polymer 3 (amorphous polymer) (VMA%=0, Mw=57300)	insoluble, non-uniform	insoluble, non-uniform	insoluble, non-uniform
Polymer 4 (amorphous polymer) (VMA%=0, Mw=31400)	barely soluble, non-uniform	barely soluble, non-uniform	barely soluble, non-uniform
Polymer 5 (amorphous polymer) (VMA%=0, Mw=22500)	slightly soluble, non-uniform	barely soluble, non-uniform	barely soluble, non-uniform
Polymer 6 (crystalline polymer) (VMA%=50, Mw=211000)	polymer swollen, uniform	polymer swollen, uniform	polymer swollen, uniform
Polymer 7 (crystalline polymer) (VMA%=50, Mw=113000)	soluble, uniform	soluble, uniform	soluble, uniform
Polymer 8 (crystalline polymer) (VMA%=50, Mw=63300)	soluble, uniform	soluble, uniform	soluble, uniform
Polymer 9 (crystalline polymer) (VMA%=50, Mw=32800)	soluble, uniform	soluble, uniform	soluble, uniform
Polymer 10 (crystalline polymer) (VMA%=50, Mw=23400)	soluble, uniform	soluble, uniform	soluble, uniform
Polymer 11 (crystalline polymer) (VMA%=50, Mw=13000)	soluble, uniform	soluble, uniform	soluble, uniform
Polymer 17 (crystalline polymer) (VMA%=30, Mw=34000)	soluble, uniform	soluble, uniform	soluble, uniform
Polymer 12 (crystalline polymer) (VMA%=80, Mw=29600)	soluble, uniform	soluble, uniform	soluble, uniform

The amorphous polymers obtained using BMA and MMA as the monomers, even with low molecular weights (for example, Mw 30,000 or lower) dissolve only barely or not at all in polypropylene glycol, but the crystalline polymers prepared from monomers containing VMA as the crystalline monomer in addition to BMA and MMA dissolve readily in polypropylene glycol even with high molecular weights (for example, 100,000 or higher). Since polypropylene glycol is very commonly used to produce polyurethanes, the use of crystalline polymers is very advantageous compared to amorphous polymers for producing polyurethanes.

10 Example 7

The example which follows concerns production of moisture-curable hot-melt adhesives, a method of evaluating them, and the results of the evaluation. The melt viscosities of the moisture-curable hot-melt adhesives were measured using a BROOKFIELD melt viscometer. The spindle used was a #27, and the measuring temperature was 121°C.

The initial strengths of the moisture-curable hot-melt adhesives were measured using a pendulum-type impact tester described in JIS K 6855 (which corresponds to ASTM D 950). The adhesive was coated on the 12.5 x 25 mm surface of a 12.5 x 25 x 10 mm birch wood material (coating temperature: 120°C), immediately after which it was sprinkled with glass beads with sizes from 0.08-0.13 mm and adhered to a 12.5 x 45 x 45 mm birch wood material. The test piece was compressed by hand, and the squeezed adhesive was removed. The impact strength was measured 1 minute after adhesion. The data was recorded as the average of measurement results for 5 test pieces.

25 The shear adhesive strength was measured according to JIS K 6851. The adhesive was applied at both ends of a 2.5 x 10 x 0.5 cm birch wood panel at 120°C, immediately after which it was sprinkled with glass beads with sizes from 0.08-0.13 mm and adhered to another birch wood panel of the same size with an adhesion area of 3.125 cm² while compressing by hand. The adhesion test piece was conditioned at 25°C, 50% relative humidity for 7 days, and then the strength

was measured with a tensile tester at a tension rate of 50 mm/min. The data was recorded as the average of measurement results for 3 test pieces.

The adhesive samples in the following examples were prepared according to the following general preparation of moisture-curable hot-melt adhesives. That is, a polyol as the polyurethane material and a pre-prepared polymer (either amorphous or crystalline) were degassed and dried for 4 hours at 120°C under vacuum prior to use. The polyol and polymer were then transferred to a 100 ml open-top reactor while still molten, and the isocyanate component MILLIONATE MT-F (4,4'-diphenylmethane diisocyanate (MDI)) and the catalyst DMDEE were added and mixed therewith by vigorous stirring for 3 minutes at 120°C under dry nitrogen. The reactor was then placed in a vacuum oven for heating and degassing for 30 minutes at 120°C under vacuum. After the heating and degassing were complete, the contents of the reactor were poured into a metal container (tube) which was then sealed. The container was kept in a desiccator at room temperature. The contents of the container (adhesives) were heated in the container at 120°C for 2 hours prior to being used in a molten state.

Some of the polymers prepared previously were combined with the polyurethane prepolymer components by the general preparation of moisture-curable hot-melt adhesives described above in the proportions listed in Table 7, to make moisture-curable hot-melt adhesives. Since the amorphous polymers did not dissolve in polyols as demonstrated in Example 6, the adhesives were prepared by the following procedure.

1) A polyurethane prepolymer comprising 58.68 parts by weight of RUCOFLEX S105-30, 25.15 parts by weight of SANNIX PP1000, 16.08 parts by weight of MILLIONATE MT-F and 0.10 part by weight of DMDEE was prepared based on the above-mentioned general preparation of moisture-curable hot-melt adhesives. (An amorphous polymer was not added.) The polyurethane prepolymer prepared in this manner will be referred to as prepolymer A.

2) Thirty grams of polymer 4 (amorphous polymer) was dissolved in MEK (solid portion: 48.8%). The solution will be referred to as solution B.

3) Twenty grams of prepolymer A and 40.98 grams of solution B were combined and mixed by stirring to uniformity at room temperature.

4) The resulting mixture was placed in a vacuum oven and the MEK was totally removed at 120°C under vacuum. The resulting final product will be referred to as adhesive C1. This adhesive was used for comparison.

The compositions of the adhesives prepared in this manner are listed in Table 7. The results of measuring the melt viscosities, initial impact strengths and overlap shear adhesive strengths of the adhesives are also summarized in Table 7. Adhesives 1-9 are the adhesives according to the invention.

Table 7

	Adhesive									
	CI	1	2	3	4	5	6	7	8	9
Polymer 4 (amorphous polymer) (VMA=0%, Mw=31400)	50.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Polymer 9 (crystalline polymer) (VMA=50%, Mw=32800)	0.00	50.00	0.00	0.00	0.00	0.00	0.00	0.00	10.00	80.00
Polymer 17 (crystalline polymer) (VMA=30%, Mw=34000)	0.00	0.00	50.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Polymer 8 (crystalline polymer) (VMA=50%, Mw=63300)	0.00	0.00	0.00	50.00	0.00	0.00	0.00	0.00	0.00	0.00
Polymer 20 (crystalline polymer*) (VMA=50%, Mw=33400)	0.00	0.00	0.00	0.00	50.00	0.00	0.00	0.00	0.00	0.00
Polymer 23 (crystalline polymer*) (VMA=50%, Mw=50700)	0.00	0.00	0.00	0.00	0.00	50.00	0.00	0.00	0.00	0.00
Polymer 12 (crystalline polymer) (VMA=80%, Mw=29600)	0.00	0.00	0.00	0.00	0.00	0.00	50.00	0.00	0.00	0.00
Polymer 11 (crystalline polymer) (VMA=50%, Mw=1300)	0.00	0.00	0.00	0.00	0.00	0.00	0.00	50.00	0.00	0.00
RUCOFLEX SI05-30	29.31	29.31	29.31	29.31	29.31	29.31	29.31	29.31	52.80	11.69
SANNIX PP1000	12.56	12.56	12.56	12.56	12.56	12.56	12.56	12.56	22.63	5.01
MILLIONATE MT-F	8.03	8.03	8.03	8.03	8.03	8.03	8.03	8.03	14.47	3.20
DMDEE	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10
NCO/OH ratio	1.6	1.6	1.6	1.6	1.6	1.6	1.6	1.6	1.6	1.6
Polymer/polyurethane prepolymer ratio	50/50	50/50	50/50	50/50	50/50	50/50	50/50	50/50	10/90	80/20
Melt viscosity (cP) 121°C	179000	9350	45100	25400	6910	3340	2160	3500	9410	7260
Initial impact strength (kg·cm)	3.5	15.5	10.5	15.6	14.5	23.0	5.1	21.8	18.0	29.2
Overlap shear adhesive strength (kg/cm ²)	28.0	33.9	43.7	41.6	44.0	40.3	15.2	36.0	78.7	33.3

* Crystalline polymer containing alkyl acrylamide

The mixture of the polyurethane prepolymer and amorphous polymer (adhesive C1) had a very high melt viscosity. Such a viscosity is not suitable for application of adhesives with common applicators. Adhesive C1 also had
5 inadequate initial impact strength. In contrast, the mixtures of the polyurethane prepolymer and crystalline polymers (adhesives 1-9) had both high initial impact strength and shear adhesive strength, with low melt viscosities. Of adhesives 1-9 according to the invention, those containing crystalline polymers comprising alkyl acrylamides (adhesives 4 and 5) had very high initial impact strength and shear
10 adhesive strength, with very low melt viscosities.

Example 8

In this example, moisture-curable hot-melt adhesives were prepared using polymers made by polymerization in polypropylene glycol.

15 A 62.56 part by weight portion of the polymer polymerized in polypropylene glycol (polymers 24-27 described in Example 5) (of which 50 parts by weight was the polymer and the remaining 12.56 parts by weight was polypropylene glycol) and 29.31 parts by weight of RUCOFLEX S105-30 were placed in an open-top reactor, and heated to dryness for 4 hours at 120°C under vacuum in a vacuum oven. After
20 next adding 8.03 parts by weight of MILLIONATE MT-F and 0.10 part by weight of DMDEE, the mixture was stirred for 3 minutes at 120°C under nitrogen. The reaction was again placed in the vacuum oven and heated for 30 minutes at 120°C under vacuum for heating and degassing. The contents of the reactor were poured into a metal container (tube) which was then sealed. The container was kept in a
25 desiccator at room temperature. The contents of the container (adhesive) were heated for two hours in the container for complete reaction prior to use.

The compositions and properties of the adhesives prepared in this manner are listed in Tables 8 and 9. Adhesive C2 is a comparative adhesive using polymer 24 (amorphous polymer), and adhesives 10-12 are adhesives according to the
30 invention.

Table 8

	Adhesive C2	Adhesive 10	Adhesive 11	Adhesive 12
Polymer 24 (amorphous polymer + PPG*)	62.56	0.00	0.00	0.00
Polymer 25 (crystalline polymer + PPG*)	0.00	62.56	0.00	0.00
Polymer 26 (crystalline polymer + PPG*)	0.00	0.00	62.56	0.00
Polymer 27 (crystalline polymer** + PPG*)	0.00	0.00	0.00	62.56
RUCOFLEX S105-30	29.31	29.31	29.31	29.31
MILLIONATE MT-F	8.03	8.03	8.03	8.03
DMDEE	0.10	0.10	0.10	0.10
NCO/OH ratio	1.6	1.6	1.6	1.6
Polymer/polyurethane prepolymer ratio	50/50	50/50	50/50	50/50

* Polypropylene glycol

** Crystalline polymer containing alkyl acrylamide

Table 9

	Adhesive C2	Adhesive 10	Adhesive 11	Adhesive 12
BMA/MMA polymer (amorphous polymer) VMA=0%, Mw=37100	50.00	0.00	0.00	0.00
VMA/BMA/MMA polymer (amorphous polymer) VMA=50%, Mw=37100	0.00	50.00	0.00	0.00
VMA/BMA/MMA polymer (amorphous polymer) VMA=30%, Mw=37300	0.00	0.00	50.00	0.00
VMA/IOA/DMAA polymer (amorphous polymer) VMA=50%, Mw=53200	0.00	0.00	0.00	50.00
RUCOFLEX S105-30	29.31	29.31	29.31	29.31
SANNIX PP1000	12.56	12.56	12.56	12.56
MILLIONATE MT-F	8.03	8.03	8.03	8.03
DMDEE	0.10	0.10	0.10	0.10
NCO/OH ratio	1.6	1.6	1.6	1.6
Polymer/polyurethane prepolymer ratio	50/50	50/50	50/50	50/50
Melt viscosity (cP) 121°C	>500000*	16080	46000	15950
Initial impact strength (kg·cm)	2.2	21.5	8.0	20.0
Overlap shear adhesive strength (kg/cm ²)	28.8	34.4	42.9	60.0

*Beyond the maximum measurable value of the melt viscometer

- The mixture of the polyurethane prepolymer and amorphous polymer
- 5 (adhesive C2) had low initial impact strength and a very high melt viscosity. Such a high viscosity results in extreme difficulty in applying the hot-melt adhesive to adherends. In contrast, the mixtures of the polyurethane prepolymer and crystalline polymers (adhesives 10-12) had both high initial impact strength and shear adhesive strength, with low melt viscosities. Of the adhesives according to the invention, the
- 10 one containing a crystalline polymer comprising an alkyl acrylamide (adhesive 12) had very high initial impact strength and overlap shear adhesive strength, with a very low melt viscosity.

Claims

1. A moisture-curable hot-melt composition comprising a polyurethane prepolymer and a crystalline polymer obtained from monomer(s) containing at least one crystalline ethylenically unsaturated monomer with no active hydrogen.
- 5
2. The composition according to claim 1, wherein said crystalline polymer is a polymer obtained from monomer(s) containing a crystalline ethylenically unsaturated compound with no active hydrogen and an amorphous ethylenically unsaturated compound with no active hydrogen.
- 10
3. The composition according to claim 1, wherein said crystalline polymer has a melting point of 15-135°C.
4. The composition according to claim 2, wherein said crystalline
- 15 polymer has a melting point of 15-135°C.
5. The composition according to claim 3, wherein said crystalline monomer is a (meth)acrylate ester of a non-tertiary alcohol, the alkyl portion of which comprises 16 or more carbon atoms.
- 20
6. The composition according to claim 4, wherein said crystalline monomer is a (meth)acrylate ester of a non-tertiary alcohol, the alkyl portion of which comprises 16 or more carbon atoms.
- 25
7. The composition according to claim 2, wherein said amorphous monomer is selected from the group consisting of (meth)acrylate esters of non-tertiary alcohols, the alkyl portions of which comprise no more than 15 carbon atoms, (meth)acrylate esters of alcohols other than alkyl alcohols, vinyl esters, vinyl ethers, alkyl acrylamides, styrene, acrylonitrile, N-vinyl lactams, vinylidene chloride
- 30 and vinyltoluene.

8. The composition according to claim 2, wherein said crystalline monomer contains an alkyl acrylamide.

9. The composition according to claim 1, wherein said monomers
5 include 5-100 wt% of a crystalline monomer.

10. The composition according to claim 1, wherein said monomers include 10-90 wt% of a crystalline monomer.

10 11. The composition according to claim 1, wherein said monomers include 30-80 wt% of a crystalline monomer.

12. The composition according to claim 1, wherein the weight average molecular weight of said crystalline polymer is between 5,000 and 100,000.
15

13. The composition according to claim 1, wherein the weight average molecular weight of said crystalline polymer is greater than 30,000 and up to 100,000.

20 14. The composition according to claim 1, wherein the weight average molecular weight of said crystalline polymer is greater than 30,000 and up to 60,000.

15. The composition according to claim 1 which contains said crystalline
25 polymer at an amount of 5-90 wt%.

16. The composition according to claim 1 which contains said crystalline polymer at an amount of 5-80 wt%.

30 17. The composition according to claim 1 which contains said crystalline polymer at an amount of 5-70 wt%.

18. The composition according to claim 1 which has a melt viscosity at 121°C (250°F) of 50,000 centipoise or lower.

5 19. The composition according to claim 1 which has an initial impact strength of 5 kg·cm or greater.

20. The composition according to claim 1 which has a final strength of 15 kg/cm² or greater.

10

INTERNATIONAL SEARCH REPORT

Int. .tional Application No

PCT/US 97/18391

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 C09J175/04

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 C09J

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	WO 96 35734 A (MINNESOTA MINING & MFG) 14 November 1996 see claims 1-16 ---	1-20
A	WO 94 24219 A (LANDEC CORP) 27 October 1994 see page 8, line 19 - page 10, line 4; claim 1 ---	1
A	DATABASE WPI Section Ch, Week 9223 Derwent Publications Ltd., London, GB; Class A14, AN 92-187565 XP002059798 & JP 04 114 080 A (SEKISUI CHEM IND CO LTD), 15 April 1992 see abstract --- -/--	1

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "&" document member of the same patent family

Date of the actual completion of the international search

20 March 1998

Date of mailing of the international search report

31/03/1998

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Goovaerts, R

INTERNATIONAL SEARCH REPORT

Int. l. Application No
PCT/US 97/18391

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>US 5 472 785 A (TANGEN JOHN C ET AL) 5 December 1995 see column 8, line 64 - column 9, line 30; claim 1</p> <p style="text-align: center;">-----</p>	1

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 97/18391

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9635734 A	14-11-96	AU 5430096 A EP 0824558 A	29-11-96 25-02-98
WO 9424219 A	27-10-94	US 5665822 A CA 2159476 A EP 0695331 A JP 9500662 T	09-09-97 27-10-94 07-02-96 21-01-97
US 5472785 A	05-12-95	AU 681666 B AU 6805294 A BR 9408561 A CA 2187524 A EP 0755414 A WO 9527748 A JP 10500708 T	04-09-97 30-10-95 19-08-97 19-10-95 29-01-97 19-10-95 20-01-98

Form PCT/ISA/210 (patent family annex) (July 1992)

THIS PAGE BLANK (USPTO)